

South Dakota Board of Pharmacy

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WHAT: Board Policy Statement # 17-06-02

WHY: WHAT INFORMATION MAY BE MODIFIED ON A WRITTEN OR E-PRESCRIBED CONTROLLED

SUBSTANCE (CS) PRESCRIPTION

INTRODUCTION

Understanding what may or may not be changed on a CS prescription is one of the many ways to continue to combat drug diversion while helping patients obtain their medications expeditiously. The rules regarding what can be modified apply to both written prescriptions and electronically prescribed prescriptions. The following summarizes changes that may be made to a prescription for a CII – CV without consulting or after consulting with the prescribing practitioner and what changes may never be made.

GUIDELINES

A pharmacist may never modify:

- 1. Patient name
- 2. Name of controlled substance (except where generic substitution permitted)
- 3. **Signature** of the practitioner

The following may be added or modified without consulting the practitioner if information can be reliably obtained:

- 1. Patient address
- 2. Practitioner address
- 3. Practitioner telephone number
- 4. ¹Quantity may be modified <u>ONLY</u> in conjunction with change of **Strength.** The total quantity dispensed cannot exceed the total dosage initially authorized.

Example: A prescription is written for methylphenidate HCl with directions 5mg (5 ml) by mouth twice daily and a quantity of 300 mL to be dispensed. The pharmacy carries 10 mg/5ml strength. The pharmacy **may** fill the prescription using the 10 mg/5ml methylphenidate HCl and change the dose and quantity accordingly. In this example, the pharmacist may change the dose to 2.5 ml (twice daily) and the quantity dispensed to 150 ml.

- ✓ The pharmacist must <u>document</u> the new quantity, strength, date, and pharmacist initials on the prescription
- 5. **Practitioner DEA number** may be **added**. However, <u>do not add a DEA number when the legitimacy of the prescription (i.e., prescriber or DEA number) is in question</u>. Only add the DEA number when it can be obtained from a validated source.

The following may be added or modified after consulting with a practitioner (may not be an agent of the practitioner). The pharmacist should document all consultations and note any changes on the prescription.

1. **Date of issue** may be added <u>but not changed.</u> A pharmacist may **not change** a "do not fill until date" even if the provider is consulted. A pharmacist <u>may</u> fill prior to a "do not fill until" date in extenuating circumstances <u>and</u> after consulting the provider.

Example: A prescription bears a "do not fill until 3/29" notation. Today's date is 3/26. The patient is leaving for a two-week vacation and requests that it be filled. After obtaining approval by the provider, you may fill the prescription.

- ✓ The pharmacist must <u>document</u> the date, reason for early fill, "prescriber consulted," and pharmacist initials on the prescription
- 2. Drug Quantity and Strength unless it falls under the example previously discussed¹
 - Includes situations where the acetaminophen strength is missing or incorrect in hydrocodone combination products. The prescriber should be contacted to verify strength of acetaminophen.
- 3. **Directions** for use unless it falls under the example previously discussed¹
- 4. **Dosage form** (capsules and tablets are not interchangeable)
- 5. Refill instructions for controlled substances III-IV
- 6. Practitioners **printed** name (NOT practitioner's signature)

Remember, a pharmacist is expected to use their professional judgment and knowledge to determine when It is appropriate to make changes to any prescription including a prescription for a controlled substance.

BOARD APPROVAL/ADOPTION: June 17, 2017